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Substitute for form 1449A/B/PTO		<b>Complete if Known</b>	
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (Use as many sheets as necessary)		Application Number	10/572,958
		Filing Date	March 21, 2006
		First Named Inventor	Dizhong Chen
		Art Unit	1114
		Examiner Name	WET YET Assigned STOCKTON
Sheet 1 of 3	Attorney Docket Number	14090-00003-US1	

U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. <sup>1</sup>	Document Number	Publication Date	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code <sup>2</sup> (if known)	MM-DD-YYYY		
	1	US 6,552,065	04-22-03	Remiszewski et al.	

FOREIGN PATENT DOCUMENTS							
Examiner Initials*	Cite No. <sup>1</sup>	Foreign Patent Document		Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	1 <sup>6</sup>
		Country Code <sup>3</sup>	Number <sup>4</sup> -Kind Code <sup>5</sup> (if known)				
	2	WO	2000/042022	07-20-2000	Warner-Lambert Co. et al.		
	3	WO	2003/077855	09-25-2003	Array Biopharma, Inc. et al.		
	4	WO	2003/077914	09-25-2003	Array Biopharma, Inc. et al.		
	5	WO	2003/087089	10-23-2003	Teijin Limited et al.		
	6	WO	2003/000682	01-03-2003	Merck & Co., Inc. et al.		
	7	WO	2003/000254	01-03-2003	Japan Tobacco Inc. et al.		
	8	WO	2002/050062	06-27-2002	Neurogen Corporation et al.		
	9	WO	2001/047883	07-05-2001	Japan Tobacco Inc. et al.		
	10	WO	2001/005390	01-25-2001	Warner-Lambert Co. et al.		
	11	WO	2001/012604	02-22-2001	Aventis Cropscience GmbH et al.		
	12	WO	2001/005393	01-25-2001	Warner-Lambert Co. et al.		
	13	WO	2001/000207	01-04-2001	Merck & Co., Inc. et al.		
	14	WO	2001/000213	01-04-2001	Merck & Co., Inc. et al.		
	15	WO	2003/066579	08-14-03	AXYS Pharmaceuticals		
	16	WO	2001/38322	05-31-01	Methylgene, Inc.		

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	17	Wade, P.A. "Transcriptional control at regulatory checkpoints by histone deacetylases: molecular connections between cancer and chromatin" Hum. Mol. Genet. Vol. 10, No. 7, pgs. 693-698 (2001)		
	18	De Ruijter, A.J.M. et al, "Histone deacetylases (HDACs): characterization of the classical HDAC family" Biochem. J., 370, pgs. 737-749 (2003)		
	19	Richon, V.M. et al, "Second generation hybrid polar compounds are potent inducers of transformed cell differentiation" Proc. Natl. Acad. Sci. USA, Vol. 93, pgs. 5705-5708 (1996)		
	20	Richon, V.M. et al, "A class of hybrid polar inducers of transformed cell differentiation inhibits histone deacetylases" Proc. Natl. Acad. Sci. USA, Vol. 95, pgs. 3003-3007 (1998)		

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Examiner Signature	/Laura L. Stockton/	Date Considered	12/16/2008
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	21	Butler, L.M. et al, "Suberoylanilide hydroxamic acid, an inhibitor of histone deacetylase, suppresses the growth of prostate cancer cells in vitro and in vivo", Cancer Res. 60, 5165-5170 (2000)		
	22	Yoshida, M. et al, "Potent and Specific Inhibition of Mammalian Histone Deacetylase Both <i>in Vivo</i> and <i>in Vitro</i> by Trichostatin A" J. Biol. Chem., Vol. 265, No. 28, pgs. 17174-17179 (1990)		
	23	Kijima, M. et al, "Trapoxin, an Antitumor Cyclic Tetrapeptide, Is an Irreversible Inhibitor of Mammalian Histone Deacetylase" J. Biol. Chem., Vol. 268, No. 30, pgs. 22429-22435 (1993)		
	24	Bouchain, G. et al, "Development of Potential Antitumor Agents. Synthesis and Biological Evaluation of a New Set of Sulfonamide Derivatives as Histone Deacetylase Inhibitors" J. Med. Chem., 46, 820-830 (2003)		
	25	Steffan, J.S. et al, "Histone deacetylase inhibitors arrest polyglutamine-dependent neurodegeneration in <i>Drosophila</i> " Nature, Vol. 413, pgs. 739-743, 18 October, 2001		
	26	Schindler, G. et al, "Dissociation between Interleukin -18 mRNA and Protein Synthesis in Human Peripheral Blood Mononuclear Cells" J. Biol. Chem., Vol. 265, No. 18, pgs. 10232-10237 (1990).		
	27	Carballo et al, "Feedback Inhibition of Macrophage Tumor Necrosis Factor- $\alpha$ Production by Tristetraprolin" Science, 1998, Vol. 281, pgs. 1001-1005		
	28	Dinarello, C.A. and Moldawer L.L. "Proinflammatory and anti-inflammatory cytokines in rheumatoid arthritis. A primer for clinicians." 3 <sup>rd</sup> Edition, Amgen Inc., 2002.		
	29	A. Inoue and D. Fujimoto, "Enzymatic Deacetylation of Histone" Biochemical Biophysical Research Communications, 1969, Vol. 36, No. 1, pgs. 146-150		
	30	J. Taunton et al, "A Mammalian Histone Deacetylase Related to the Yeast Transcriptional Regulator Rpd3p" Science, April 19, 1996, Vol. 272, pgs. 408-411		
	31	P.A. Wade et al, "Purification of a Histone Deacetylase Complex from <i>Xenopus Laevis</i> : Preparation of Substrates and Assay Procedures" Methods in Enzymology, 1999, Vol. 304, pgs. 715-725		
	32	A. Ito et al, "p300/CBP-mediated p53 acetylation is commonly induced by p53-activating agents and inhibited by MDM2" EMBO Journal 2001, Vol. 20, No. 6, pgs. 1331-1340		
	33	B.D. Strahl and C.D. Allis, "The Language of Covalent Histone Modifications" Nature, January 6, 2000, Vol. 403, pgs. 41-45		
	34	B. Helweg and M. Jung, "A Microplate Reader-Based Nonisotopic Histone Deacetylase Activity Assay" Anal. Biochem. 2002, Vol. 302, pgs. 175-183		

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	35	Ito et al, "A Molecular Mechanism of Action of Theophylline: Induction of Histone Deacetylase Activity to Decrease Inflammatory Gene Expression" Proc. Natl. Acad. Sci. USA 2002, Vol. 99, No. 13, pgs. 8921-8926		
	36	K.J. Bitterman et al, "Inhibition of Silencing and Accelerated Aging by Nicotinamide, a Putative Negative Regulator of Yeast Sir2 and Human SIRT1" J. Biol. Chem. 2002, Vol. 277, No. 47, pgs. 45099-45107		
	37	S. Milutinovic et al, "Proliferating Cell Nuclear Antigen Associates with Histone Deacetylase Activity, Integrating DNA Replication and Chromatin Modification" J. Biol. Chem. 2002, Vol. 277, No. 23, pgs. 20974-20978		
	38	Still et al, "Rapid Chromatographic Technique for Preparative Separations with Moderate Resolution" J. Org. Chem., Vol. 43, No. 14, pgs. 2923-2925 (1978)		
	39	Remiszewski et al, "Inhibitors of Human Histone Deacetylase: Synthesis and Enzyme and Cellular Activity of Straight Chain Hydroxamates" J. Med. Chem., 2002, Vol. 45, No. 4, pgs. 753-757		
	40	Baudy et al, "Design, Synthesis, SAR, and Biological Evaluation of Highly Potent Benzimidazole-Spaced Phosphono- $\alpha$ -Amino Acid Competitive NMDA Antagonists of the AP-6 Type" J. Med. Chem. 2001, 44, 1516-1529		
	41	Yu-Hua Ji et al, "Tris-benzimidazole derivatives: design, synthesis and DNA sequence recognition" Bioorganic & Medical Chemistry 9, pp 2905-2919 (2001) et al.		

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